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MONOCRYL™

ar
bg КОНЕЦ
cs ŠICÍ MATERIÁL
da SUTUR
de NAHTMATERIAL
el PAMMA
en SUTURE
es SUTURA
et ÕMBLUSMATERJAL
fi OMMELAINE
fr FIL DE SUTURE
hu VARRÓANYAG
it SUTURA
kk ТІГУ МАТЕРИАЛЫ
ko 봉합사
lt SIŪLAS
lv ĶIRURĢISKAIS DIEGS
zh-cn 缝线
nl HECHTMATERIAAL
zh-tw 縫合線
no SUTUR
pl NICI CHIRURGICZNE
pt FIO DE SUTURA
ro FIR DE SUTURĂ
ru ШОВНЫЙ МАТЕРИАЛ
sk CHIRURGICKÁ NIŤ
sv SUTUR
tr SÜTÜR

خيط
جر

La résorption est pratiquement complète au bout de 91 à 119 jours.

CONTRE-INDICATIONS
Comme il est résorbable, ce fil de suture (coloré et incolore) ne doit pas être utilisé lorsqu'un rapprochement prolongé des tissus soumis à un stress est nécessaire.
Le fil de suture incolore MONOCRYL™, en particulier, ne doit pas être utilisé pour la fermeture des tissus fasciaux.

AVERTISSEMENTS
L'innocuité et l'efficacité du fil de suture MONOCRYL™ n'ont pas été établies dans les cas suivants : tissus nerveux, tissus cardiovasculaires, microchirurgie et chirurgie ophtalmique.
L'utilisateur doit connaître les procédures et les techniques chirurgicales relatives à l'utilisation de fils de suture résorbables avant d'utiliser le fil de suture MONOCRYL™ pour la fermeture d'une plaie ainsi que le risque de déhiscence de la plaie qui dépend du site opératoire et de la nature du fil de suture. Le choix du fil de suture par les médecins est fonction de ses propriétés *in vivo* (voir la rubrique MODE D'ACTION).
Il convient de respecter les procédures chirurgicales habituelles nécessaires au contrôle de la contamination ou de l'infection de la plaie.
Comme ce fil de suture est résorbable, le chirurgien peut envisager d'utiliser en complément des fils de suture non résorbables pour la fermeture de plaies susceptibles de subir une expansion, un étirement ou une distension, ou exigeant un suction supplémentaire.
Ne pas resteriliser/réutiliser. La réutilisation de ce dispositif (ou de parties de ce dispositif) peut créer un risque de dégradation du produit, susceptible d'entraîner une défaillance du dispositif et/ou une contamination croisée, ce qui peut provoquer une infection ou la transmission d'agents pathogènes transmissibles par le sang aux patients et aux utilisateurs.
Dans la mesure où le fil de suture est un matériau résorbable, il peut se comporter transitoirement comme un corps étranger. Comme tout corps étranger, ce produit peut aggraver une infection existante.

PRÉCAUTIONS
Une attention particulière doit être portée à la manipulation de ce fil et de tout autre matériel de suture afin de ne pas risquer de les détériorer. Ne pas écraser ou pincer le fil lors de l'application d'instruments chirurgicaux tels que des pinces ou des porte-aiguilles.
Les fils de suture cutanés restant en place pendant plus de 7 jours peuvent entraîner une irritation localisée et doivent être coupés ou retirés selon les indications.
Dans certaines circonstances, notamment lors de procédures orthopédiques, une immobilisation des articulations par soutien externe peut être utilisée à la discrétion du chirurgien.
Il convient d'être prudent en cas d'utilisation de fils de suture résorbables au niveau de tissus mal irrigués, car il existe alors un risque de rejet de la suture et de retard de la résorption.

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For recognized legal manufacturer, refer to product label.

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Instructions for use

en

MONOCRYL™

Poliglecaprone 25 (MONOFILAMENT)
STERILE SYNTHETIC ABSORBABLE SURGICAL SUTURE

DESCRIPTION
MONOCRYL™ Suture is a sterile, monofilament, synthetic, absorbable surgical suture composed of a copolymer of glycolide and (epsilon) ε-caprolactone. The empirical molecular formula of the polymer is $(C_2H_2O_2)_m$ $(C_6H_{10}O_2)_n$.
Poliglecaprone 25 copolymer has been found to be nonantigenic, nonpyrogenic and elicits only a slight tissue reaction during absorption.
MONOCRYL™ Suture is available undyed and dyed with D&C Violet No. 2 (Color Index 60725) to enhance visibility in the surgical field.
MONOCRYL™ Suture is available in a range of gauge sizes and lengths, non-needled or attached to needles of varying types and sizes as described in the HOW SUPPLIED section.
MONOCRYL™ Suture complies with the requirements of the European Pharmacopoeia (Ph. Eur.) for Sterile Synthetic Absorbable Monofilament Sutures and United States Pharmacopeia (USP) for Absorbable Surgical Sutures except for a slight oversize in diameter for all gauges as shown in the table below.

USP Suture Size	Metric Suture Size	Maximum Oversize
6-0	0.7	0.049 mm
5-0	1	0.033 mm
4-0	1.5	0.045 mm
3-0	2	0.067 mm
2-0	3	0.055 mm
0	3.5	0.088 mm
1	4	0.066 mm

The European Pharmacopoeia recognizes units of measure Metric and Ph. Eur. Sizes as equivalent, which is reflected on the labeling.

INDICATIONS
MONOCRYL™ Sutures are indicated for use in general soft tissue approximation and/or ligation, but not for use in cardiovascular or neurological tissues, microsurgery or ophthalmic surgery.

APPLICATIONS
Sutures should be selected and implanted depending on patient condition, surgical experience, surgical technique and wound size.

PERFORMANCE / ACTIONS
MONOCRYL™ Suture elicits a minimal initial inflammatory reaction in tissues and ingrowth of fibrous connective tissue. Progressive loss of tensile strength and eventual absorption occurs by means of hydrolysis, where the polymer degrades to adipic acid, which is subsequently absorbed and metabolized in the body. Absorption begins as loss of tensile strength followed by a loss of mass. Implantation studies in rats show the following profile:

DYED		UNDYED	
Post Implantation	Approx. Original Strength Remaining	Post Implantation	Approx. Original Strength Remaining
7 days	60% to 70%	7 days	50% to 60%
14 days	30% to 40%	14 days	20% to 30%
All of the original tensile strength is essentially lost by 28 days post-implantation.		All of the original tensile strength is essentially lost by 21 days post-implantation.	

Absorption is essentially complete between 91 and 119 days.

CONTRAINDICATIONS
This suture (dyed and undyed), being absorbable, should not be used where extended approximation of tissues under stress is required.
Undyed MONOCRYL™ Sutures, in particular, should not be used to close fascial tissue.

WARNINGS
The safety and effectiveness of MONOCRYL™ Sutures has not been established in the following areas – neural tissue, cardiovascular tissue, microsurgery and ophthalmic surgery.
Users should be familiar with surgical procedures and techniques involving absorbable sutures before employing MONOCRYL™ Suture for wound closure, as risk of wound dehiscence may vary with the site of application and the suture material used. Medical professionals should consider the *in vivo* performance (under PERFORMANCE/ACTIONS section) when selecting a suture.
Acceptable surgical practice should be followed for the management of contaminated or infected wounds.
As this is an absorbable suture material, the use of supplemental nonabsorbable sutures should be considered by the surgeon in the closure of sites which may undergo expansion, stretching or distension, or that may require additional support.
Do not resterilize/reuse. Reuse of this device (or portions of this device) may create a risk of product degradation, which may result in device failure and/or cross-contamination, which may lead to infection or transmission of blood-borne pathogens to patients and users.
As an absorbable suture, this suture may act transiently as a foreign body. Like all foreign bodies, this product may potentiate infection.

PRECAUTIONS
In handling this or any other suture material, care should be taken to avoid damage. Avoid crushing or crimping damage due to application of surgical instruments such as forceps or needle holders.
Skin sutures which must remain in place longer than 7 days may cause localized irritation and should be snipped off or removed as indicated.
Under some circumstances, notably orthopedic procedures, immobilization of joints by external support may be employed at the discretion of the surgeon.
Consideration should be taken in the use of absorbable sutures in tissues with poor blood supply as suture extrusion and delayed absorption may occur.
Subcuticular sutures should be placed as deeply as possible to minimize the erythema and induration normally associated with the absorption process.
This suture may be inappropriate in elderly, malnourished or debilitated patients, or in patients suffering from conditions that may delay wound healing.
As with any suture material, adequate knot security requires the standard surgical technique of flat, square ties with additional throws as warranted by surgical circumstance and the experience of the surgeon. The use of additional throws may be particularly appropriate when knotting monofilament sutures.
Care should be taken to avoid damage when handling surgical needles. Grasp the needle in an area one-third (1/3) to one-half (1/2) of the distance from the attachment end to the point. Grasping in the point area could impair the penetration performance and cause fracture of the needle. Grasping at the attachment end could cause bending or breakage. Reshaping needles may cause them to lose strength and be less resistant to bending and breaking.
Users should exercise caution when handling surgical needles to avoid inadvertent needle stick injury. Broken needles may result in extended or additional surgeries or residual foreign bodies. Inadvertent needle sticks with contaminated surgical needles may result in the transmission of blood-borne pathogens. Discard used needles in "sharps" containers.

ADVERSE REACTIONS
Adverse reactions associated with the use of this device include wound dehiscence, calculi formation in urinary or biliary tracts when prolonged contact with salt solutions such as urine or bile occurs, transient inflammatory tissue reaction, and transient local irritation at the wound site.

STERILITY
MONOCRYL™ Suture is sterilized by ethylene oxide gas. Do not resterilize. Do not use if package is opened or damaged. Discard opened, unused sutures.

STORAGE
No special storage conditions required. Do not use after expiry date.

HOW SUPPLIED
Please note that not all sizes are available in all markets. Please contact your local sales representative for size availability.
MONOCRYL™ Suture is available as sterile, monofilament strands in sizes 6-0 through 1 (metric sizes 0.7 - 4), in a variety of lengths, with and without permanently attached needles.
MONOCRYL™ Suture is also available in sizes 3-0 through 1 (metric sizes 2 - 4) as sterile strands attached to CONTROL RELEASE™ removable needles, which enable the needles to be pulled off instead of being cut off.
MONOCRYL™ Suture is available in one, two or three dozen units per box.

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ar لا تُعد استخدامه / **bg**-Не използвайте повторно / **cs**-Pro jednorázové použití / **da**-Må ikke genbruges / **de**-Nicht zur Wiederverwendung / **el**-Μην επαναχρησιμοποιείτε / **en**-Do not reuse / **es**-No reutilizar / **et**-Ärge kasutage korduvalt / **fi**-Ei saa käyttää uudelleen / **fr**-Ne pas réutiliser / **hu**-Ne használja újra / **it**-Non riutilizzare / **kk**-Қайта пайдаланбаңыз / **ko**-재사용하지 마십시오 / **lt**-Tinka iki nurodytos datos / **lv**-Nelietot atkārtoti / **nl**-Niet opnieuw gebruiken / **no**-Skal ikke brukes på nytt / **pl**-Nie używać powtórnie / **pt**-Não reutilizar / **ro**-A nu se reutiliza / **ru**-Не использовать повторно / **sk**-Nepoužívajte opakovane / **sv**-Får ej återanvändas / **tr**-Tekrar kullanmayın / **zh-cn**-不得重复使用 / **zh-tw**-不得重複使用

تستخدم قبل تاريخ-**ar** / **bg**-Да се употреби до / **cs**-Datum použitelnosti / **da**-Anvendes inden / **de**-Verfallsdatum / **el**-Ημερομηνία λήξης / **en**-Use-by date / **es**-Fecha de caducidad / **et**-Viimane kasutuskuupäev / **fi**-Viimeinen käyttöpäivä / **fr**-Date de péremption / **hu**-Felhasználhatóság dátuma / **it**-Utilizzare entro il / **kk**-Жарамдылық мерзімі / **ko**-사용기한일 / **lt**-Tinka iki nurodytos datos / **lv**-Izlietot līdz (datums) / **nl**-Uiterste gebruiksdatum / **no**-Utløpsdato / **pl**-Wykorzystać do dnia / **pt**-Data de validade / **ro**-A se utiliza până la data de / **ru**-Использовать до (дата) / **sk**-Spotrebujte do / **sv**-Används före / **tr**-Son kullanma tarihi / **zh-cn**-使用有效期 / **zh-tw**-使用有效日期

LOT رمز التشغيلة-**ar** / **bg**-Партиден код / **cs**-Číslo šarže / **da**-Batchkode / **de**-Chargenbezeichnung / **el**-Κωδικός παρτίδας / **en**-Batch code / **es**-Código de lote / **et**-Partii kood / **fi**-Eräkoodi / **fr**-Code de lot / **hu**-Gyártási kód / **it**-Codice di lotto / **kk**-Партия коды / **ko**-배치 코드 / **lt**-Partijos kodas / **lv**-Partijas kods / **nl**-Batchcode / **no**-Partikode / **pl**-Kod serii / **pt**-Código de lote / **ro**-Codul lotului / **ru**-Номер партии / **sk**-Kód šarže / **sv**-Satskod / **tr**- Parti kodu / **zh-cn**-批号 / **zh-tw**-批號

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bg-CE маркировка и идентификационен номер на нотифицирания орган. Продуктът отговаря на основните изисквания на директивата за медицинските устройства 93/42/ЕИО / **cs**-Označení CE a identifikační číslo notifikované osoby. Výrobek odpovídá základním požadavkům směrnice 93/42/EHS o zdravotnických prostředcích /

STERILE EO
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EC REP ممثل معتمد لدى الاتحاد الأوروبي-**ar** / **bg**-Упълномощен представител за Европейската общност / **cs**-Autorizovaný zástupce v Evropském společenství / **da**-Autoriseret repræsentant i EU / **de**-Bevollmächtigter in der Europäischen Gemeinschaft / **el**-Εξουσιοδοτημένος αντιπρόσωπος στην Ευρωπαϊκή Κοινότητα / **en**-Authorized Representative in the European Community / **es**-Representante autorizado en la Comunidad Europea / **et**-Ametlik esindaja Euroopa Ühenduses / **fi**-Valtuutettu edustaja Euroopan yhteisössä / **fr**-Mandataire agréé dans la Communauté européenne / **hu**-Engedéllyel rendelkező képviselet az Európai Közösségben / **it**-Rappresentante autorizzato nella Comunità europea / **kk**-ЕО ресми өкілі / **ko**-유럽연합 공식 대리인 / **lt**-Įgaliotasis atstovas Europos Bendrijoje / **lv**-Pilnvarotais pārstāvis Eiropas Kopienā / **nl**-Gemachtigd vertegenwoordiger in de Europese Unie / **no**-Autorisert representant i EU / **pl**-Autoryzowany przedstawiciel we Wspólnocie Europejskiej / **pt**-Representante autorizado na Comunidade Europeia / **ro**-Reprezentant autorizat în Comunitatea Europeană /

EC REP **ru**-Официальный представитель в ЕС / **sk**-Autorizovaný zástupca pre Európske spoločenstvo / **sv**-Auktoriserad representant i Europeiska gemenskapen / **tr**-Avrupa Topluluğu Yetkili Temsilcisi / **zh-cn**-欧洲共同体授权代理 / **zh-tw**-歐洲共同體授權代表

CE 2797 **da**-CE-mærkning og identifikationsnummer for bemyndiget organ. Produktet opfylder de væsentlige krav i Direktiv 93/42/ØF om medicinsk udstyr / **de**-CE-Kennzeichnung und Kennnummer der Benannten Stelle. Das Produkt entspricht den grundlegenden Anforderungen der Richtlinie 93/42/EWG des Rates über Medizinprodukte / **el**-Σήμανση CE και Αριθμός Αναγνώρισης του Κοινοποιημένου Οργανισμού. Το προϊόν πληροί τις ουσιώδεις απαιτήσεις της Οδηγίας 93/42/ΕΟΚ περί ιατροτεχνολογικών προϊόντων / **en**-CE-mark and Identification number of Notified Body. The product meets the essential requirements of Medical Device Directive 93/42/EEC / **es**-Marca CE + número de identificación del organismo notificado. Este producto cumple los requisitos esenciales de la directiva de productos sanitarios 93/42/CEE / **et**-CE-märgis ja teavitatud asutuse registreerimisnumber. Toode vastab meditsiinseadmete direktiivi 93/42/EMÜ põhinõuetele / **fi**-CE-merkintä ja ilmoitetun laitoksen tunnusnumero. Tuote noudattaa lääkinnällisistä laitteista annetun direktiivin 93/42/ETY olennaisia vaatimuksia / **fr**-Marquage CE et numéro d'identification de l'organisme notifié. Ce produit est conforme aux exigences essentielles de la Directive sur les dispositifs médicaux 93/42/CEE / **hu**-CE-jel és a bejegyzett testület száma. A termék megfelel az orvosi készülékekről szóló 93/42/EGK számú irányelv előírásainak / **it**-Marchio CE e numero di identificazione dell'organismo notificato. Il prodotto è conforme ai requisiti essenziali della direttiva 93/42/CEE sui dispositivi medici / **kk**-CE – белгі және құзыретті органның бірегейлік нөмірі. Өнім 93/42/ЕЭК медициналық буйымдар туралы Директивасының негізгі талаптарына жауап береді / **ko**-CE마크 및 인증기관의 식별번호. 제품은 의료기기규정 93/42/EEC의 해당 요구조건을 준수합니다 / **lt**-CE ženklas ir notifikuotosios įstaigos identifikavimo numeris. Produktas atitinka pagrindinius medicininių prietaisų 93/42/EEB direktyvos reikalavimus / **lv**-CE zīme un sertifikācijas iestādes identifikācijas numurs. Izstrādājums atbilst medicīnisko ierīču direktīvā 93/42/EEK izvirzītajām pamatprasībām / **nl**-CE-markering en identificatienummer van de aangemelde instantie. Het product voldoet aan de essentiële eisen van de Richtlijn voor medische hulpmiddelen 93/42/EEG /

CE 2797 **no**-CE-merke og ID-nummer for bemyndiget organ. Produktet innfir de vesentlige krav i direktivet 93/42/EØF for medisinsk utstyr / **pl**-Znak CE i numer identyfikacyjny jednostki notyfikowanej. Produkt spełnia wymagania zasadnicze zawarte w dyrektywie 93/42/EWG dotyczącej wyrobów medycznych / **pt**-Marca CE e número de identificação do organismo notificado. O produto cumpre os principais requisitos da diretiva sobre dispositivos médicos 93/42/CEE / **ro**-Marcajul CE și numărul de identificare a organismului notificat. Acest produs corespunde cerințelor esențiale ale Directivei privind dispozitivele medicale 93/42/CEE / **ru**-Знак сертификации «CE» и идентификационный номер уведомленного органа. Изделие отвечает основным требованиям директивы о медицинском оборудовании 93/42/ЕЭС / **sk**-Označenie CE a identifikačné číslo notifikovaného orgánu. Tento produkt spĺňa základné požiadavky smernice č. 93/42/EHS o zdravotníckych pomôckach / **sv**-CE-märke och identifieringsnummer för anmält organ. Produkten uppfyller de väsentliga kraven i direktivet för medicinsk utrustning 93/42/EEG / **tr**-CE işareti ve Onaylanmış Kuruluşun kimlik numarası. Ürün, 93/42/EEC sayılı Tıbbi Cihazlar Direktifinin başlıca gerekliklerine uygundur / **zh-cn**-CE 标志和认证机构的识别号。产品符合医疗设备指令 93/42/EEC 的基本要求 / **zh-tw**-CE 標誌和認證機構的識別號碼。產品符合醫療設備指令 93/42/EEC 的基本要求

عدد الوحدات-**ar** / **bg**-Брой единици / **cs**-Počet kusů / **da**-Antal enheder / **de**-Stückzahl / **el**-Αριθμός μονάδων / **en**-Number of Units / **es**-Número de unidades / **et**-Seadmete arv / **fi**-Yksikköjen lukumäärä / **fr**-Nombre d'unités / **hu**-Az egységek száma / **it**-Numero di unità / **kk**-Бірліктер саны / **ko**-제품 개수 / **lt**-Vienetų skaičius / **lv**-Vienību skaits / **nl**-Aantal eenheden / **no**-Antall enheter / **pl**-Liczba jednostek / **pt**-Número de unidades / **ro**-Număr de unități / **ru**-Число единиц / **sk**-Počet kusov / **sv**-Antal enheter / **tr**-Ünite sayısı / **zh-cn**-件数 / **zh-tw**-數量

CR/X (X=عدد الخيوط في كل عبوة) CONTROL RELEASE™-**ar**/ **bg**-CONTROL RELEASE™ (X=Брой снопове в една опаковка) / **cs**-CONTROL RELEASE™ (X=počet vláken v jednom balení) / **da**-CONTROL RELEASE™ (X=Antal tråde pr. pakke) / **de**-CONTROL RELEASE™ (X=die Anzahl der Stränge pro Paket) / **el**-CONTROL RELEASE™ (X=Ο αριθμός των νημάτων ανά πακέτο) / **en**-CONTROL RELEASE™ (X=The number of strands per packet) / **es**-CONTROL RELEASE™ (X=número de hebras por envase) / **et**-CONTROL RELEASE™ (X=niitide arv pakendis) / **fi**-CONTROL RELEASE™ (X=lankojen määrä pakettia kohden) / **fr**-CONTROL RELEASE™ (X=nombre de brins par paquet) / **hu**-CONTROL RELEASE™ (X=A varrószálak száma csomagonként) / **it**-CONTROL RELEASE™ (X=numero di fili per confezione) / **kk**-CONTROL RELEASE™ (X=Әр орауыштағы түйіндер саны) / **ko**-CONTROL RELEASE™ (X=패킷당 가닥 수) / **lt**-CONTROL RELEASE™ (X=vijų skaičius pakuotėje) / **lv**-CONTROL RELEASE™ (X=šķiedru skaits iepakojumā) / **nl**-CONTROL RELEASE™ (X=het aantal draden per verpakking) / **no**-CONTROL RELEASE™ (X=antall tråder i hver pakke) / **pl**-CONTROL RELEASE™ (X=Liczba pasm w opakowaniu) / **pt**-CONTROL RELEASE™ (X=número de filamentos por embalagem) / **ro**-CONTROL RELEASE™ (X=numărul de filamente per pachet) / **ru**-CONTROL RELEASE™ (X=число нитей в пакете) / **sk**-CONTROL RELEASE™ (X=počet vlákien na balenie) / **sv**-CONTROL RELEASE™ (X=antalet suturer per förpackning) / **tr**-CONTROL RELEASE™ (X=Paket başına iplik sayısı) / **zh-cn**-CONTROL RELEASE™ (X=每包股线的数目) / **zh-tw**-CONTROL RELEASE™ （X=每包股數量）

MS/X (X=عدد الخيوط في كل عبوة) عديد الخيوط-**ar** / **bg**-Множество снопове (X=Брой снопове в една опаковка) / **cs**-Multistrand (X=počet vláken v jednom balení) / **da**-Multi-tråd (X=Antal tråde pr. pakke) / **de**-Mehrfachstrang (X=die Anzahl der Stränge pro Paket) / **el**-Πολλαπλά νήματα (X=Ο αριθμός των νημάτων ανά πακέτο) / **en**-Multistrand (X=The number of strands per packet) / **es**-Multihebra (X=número de hebras por envase) / **et**-Multistrand (X=niitide arv pakendis) / **fi**-Monilankainen (X=lankojen määrä pakettia kohden) / **fr**-Multibrin (X=nombre de brins par paquet) / **hu**-Többszálú (X=A szálak száma csomagonként) / **it**-Multifilo (X=numero di fili per confezione) / **kk**-Көп түйін (X=Әр орауыштағы түйіндер саны) / **ko**-다중 가닥(X=패킷당 가닥 수) / **lt**-Daugiavijis (X=vijų skaičius pakuotėje) / **lv**-Daudzšķiedru (X=šķiedru skaits iepakojumā) / **nl**-Meerdere draden (X=het aantal draden per verpakking) / **no**-Flertråded (X=antall tråder i hver pakke) / **pl**-Wielopasmowy (X=Liczba pasm w opakowaniu) / **pt**-Multifilamentos (X=número de filamentos por embalagem) / **ro**-Multifilament (X=numărul de filamente per pachet) / **ru**-Многониточный (X=число нитей в пакете) / **sk**-Viacvláknová (X=počet vlákien na balenie) / **sv**-Multistrand (X=antalet suturer per förpackning) / **tr**-Çok ipliklı (X=Paket başına iplik sayısı) / **zh-cn**-多股（X=每包股线的数目）/ **zh-tw**-多股（X=每包股數量）

افتح هنا/انزع/أزل الغلاف-**ar** / **bg**-ОТВЕПЕТЕ/ПОВДИГНЕТЕ/ОТВОРЕТЕ ТУК / **cs**-ZDE ODLOUPNĚTE/ZDVIHNĚTE/OTEVŘETE / **da**-TRÆK AF/LØFT/ÅBN HER / **de**-HIER ABZIEHEN/ANHEBEN/ÖFFNEN / **el**-ΑΠΟΚΟΛΛΗΣΤΕ/ΑΝΑΣΗΚΩΣΤΕ/ΑΝΟΙΞΤΕ ΕΔΩ / **en**-PEEL/LIFT/OPEN HERE / **es**-DESPRENDER/LEVANTAR/ABRIR AQUÍ / **et**-TÕMBA/TÕSTA/AVA SIIT / **fi**-KUORI/NOSTA/AVAA TÄSTÄ / **fr**-DÉTACHER/SOULEVER/OUVRIR ICI / **hu**-HÚZZA LE/EMELJE MEG/NYISSA FEL ITT / **it**-STRAPPARE/SOLLEVARE/APRIRE QUI / **kk**-ОСЫ ЖЕРДЕН СЫДЫРЫҢЫЗ/КӨТЕРІҢІЗ/АШЫҢЫЗ / **ko**-여기를 벗기십시오/들어 올리십시오/여십시오 / **lt**-NULUPTI/PAKELTI/ATIDARYTI ČIA / **lv**-NOPLĒST/PACELT/ATVĒRT ŠEIT / **nl**-HIER OPENTREKKEN/OMHOOGTREKKEN/OPENEN / **no**-SKRELLES AV/LØFTES/ÅPNES HER / **pl**-ODERWAĆ/UNIEŚĆ/OTWIERAĆ TUTAJ / **pt**-PUXAR/LEVANTAR/ABRIR AQUI / **ro**-DEZLIPIȚI/RIDICAȚI/DESCHIDEȚI AICI / **ru**-СНЯТЬ/ПРИПОДНЯТЬ/ОТКРЫТЬ ЗДЕСЬ / **sk**-TU ODLÚPNUŤ/NADVIHNÚŤ/OTVORIŤ / **sv**-DRA AV/LYFT/ÖPPNA HÄR / **tr**-BURADAN SOYUN/KALDIRIN/AÇIN / **zh-cn**-剥开/提起/此处开启 / **zh-tw**-剝開/提起/此處開啟

افتح هنا-**ar** / **bg**-СЪЪЛТЕ ТУК / **cs**-ZDE ODTRHNĚTE / **da**-RIV HER / **de**-HIER AUFREISSEN / **el**-ΣΚΙΣΤΕ ΕΔΩ / **en**-TEAR HERE / **es**-RASGAR AQUÍ / **et**-REBI SIIT / **fi**-REVI TÄSTÄ / **fr**-DÉCHIRER ICI / **hu**-SZAKÍTSA LE ITT / **it**-ROMPERE QUI / **kk**-ОСЫ ЖЕРДЕН ЖЫРТЫҢЫЗ / **ko**-여기를 찢으십시오 / **lt**-NUPLĖŠTI ČIA / **lv**-NOPLĒST ŠEIT / **nl**-HIER AFSCHEUREN / **no**-RIVES HER / **pl**-TU ROZERWAĆ / **pt**-RASGAR AQUI / **ro**-RUPEȚI AICI / **ru**-ОТОРВАТЬ ЗДЕСЬ / **sk**-TU ODTRHNÚŤ / **sv**-RIV HÄR / **tr**-BURADAN YIRTIN / **zh-cn**-由此撕开 / **zh-tw**-在此撕開